MAR 2 2 2004

DUPLICATE

Sheet 1 of 1

RA	FORESPTO-1449 U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE				ATTY, DOCKE	APPLICATION NO.:					
	INFORMATION DISCLOSURE STATEMENT					APPLICANT:	10/654,163				
	BY APPLICANT APPLICANT Timothy J. Guzi et al.										
	(Use several sheets if necessary)					FILING DATE: 09/03/2003	GROUP:				
	U.S. PATENT DOCUMENTS										
	'EXAMINER		DOCUMENT			ME CLASS		SUB-	SUB- FILING DATE I		
	INITIAL	L_	NUMBER		ļ			CLASS	APPRO	PRIATE	
		AA			ļ						
- 1		AB	<u> </u>	 	ļ		 		ļ		
		AC						 			
	<u></u>	AD AE	 	ļ	 		 	<u> </u>	 		
1		AF	<u> </u>		 		 	 	├		
		AG	·				 -	 	 		
·		AH					 	 	 		
		Al	 					 	 		
		AJ						 	 		
		AK			•						
	FOREIGN PATENT DOCUMENTS										
			DOCUMENT .	DATE	COUNTRY		CLASS	SUB-	TRANSI	ATION	
		L	NUMBER					CLASS	YES	NO	
		AL	DE 102 23 917 A1	12/11/2003	Germ	any					
			EP 1 199 306 A1	04/24/2002	Euro						
- 1		_	EP 0 714 898 A1	06/05/1996	Euro						
<u> </u>	$-A \nearrow$	AO	WO 02/50079 A1	06/27/2002	WIF		<u> </u>		<u> </u>		
•		AP	WP 03/091256 A1	11/06/2003	WIF		<u></u>	L	L		
۱.		OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)									
ł	Ω	AQ Adrian M. Senderowicz et al., "Preclinical and Clinical D			cal Developme	nt of Cyc	in-Depen	dent Kin	ase		
ŀ	~	AD	Modulators", J. Natl. Cancer Inst., 92(5):376-387 (March 1, 2000).								
	W	AR	Teruki Honma et al., "Structure-Based Generation of a New Class of Potent Cdk4 Inhibitors: New de Novo Design Strategy and Library Design", J. Med. Chem., 44: 4615-4627 (2001).								
ŀ	-00	AS	Translation of WO 0	3/91256 <i>A Bi</i>	sing Sun Commi	unications I td	Translatio	21 (200)	1).		
ı		AT		301200,7171	Sing Carr Commi	micanoris Liu.	T Tall Sialic	III FIOUUL	1, (1-02)		
ı		AU	-, 								
		ΑV		1	··	· · · · · · · · · · · · · · · · · · ·				$\overline{}$	
	EXAMINER		1/1	1/	DATE CO	NSIDERED	. /		-		
\downarrow		}	1. 1UC.	14		6/H	1/cc				
1	*EXAMINE	i: In	itial if reference consi	ered, Whethe	er or not citation	s in conforman	ce with M	PEP 609	: Draw I	ine	
Y	through cita applicant.	tion i	f not in conformance	and not consid	dered. Include c	opy of this form	with nex	t commu	nication t	lo	
_											

	PATE	DEC 1	5 2003 &	•				•	Shee	1 <u>1</u> of	1
FORM PTO PRINT U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE OC01620K						ET NO	T NO.: APPLICATION NO.: 10/654,163				
INFORMATION DISCLOSURE STATEMENT BY APPLICANT APPLICANT: Tim thy J. 6							Guzi	t al.	·		
(Use several sheets if necessary)						FILING DATE: 09/03/2003			GROUP/624		
				U.S. PATI	ENT DOCU						
'EXAMINE!			DOCUMENT	DATE	N/	ME	CLASS		SUB-	FILING APPRO	
INIT			NUMBER	11/05/1996	Dahi	e at at	+-	_	CLASS	MAPRO	PRIATE
	<u> </u>	AA	US 5,571,813	02/11/1997	Rühter et al.		+	+	 	 	
	4—	AB	US 5,602,136	02/11/1997			-	╫	 	 	
		AC A	US 5,602,137		Rühter et al.		+-	╂	 	┼	
<u> </u>	┼	AD	US 5,688,949	11/18/1997 01/13/1998		et al.	+-	+-	 		
		AE	US 5,707,997	07/06/1999		ey et al.	+	\vdash	 	 	
	╂-		US 5,919,815	03/21/2000		el al.	+	 	 	 	
<u></u>	-	AG	US 6,040,321	02/20/2001		et al.	+-/		+	 	
<u> </u>	/	AH	US 6,191,131 US 6,262,096	07/17/2001		et al.	++		 	 	
 	//	AJ	US 6,413,974	07/02/2002			++		 	 	
$\vdash U$		<u> </u>					-4-		4		
					ATENT DOC	UMENIS			Y 2.:.	Terre	
			DOCUMENT	DATE	COUNTRY		Cr	ASS	SUB-		LATION
			NUMBER				—		CLASS	YES	NO
		AK	EP 0 628 559	04/03/2002		rope			<u> </u>	 	
		AL	EP 1 334 973	08/13/2003	Europe		4			↓	
		АМ	WO 02/40485	05/23/2002		CT	4		<u> </u>	}	↓
1-4		AN	WO 02/50079	06/27/2002		CT			 	 	-
$\perp u$		AO	WO 95/35298					<u> </u>	┸	1	
1		OT	HER DOCUMENT	'S (Including	Author, Titl	e, Date, Pe	rtineı	nt Pa	iges, E	tc.)	
82 AP			Vesely et al., "Inhibition of Cyclin-Dependent Kinases by Purine Analogues", Eur. J. Blocham (1994), 224: 771-786.								
	AQ Kim et al., "Discovery of Aminothiazole Inhibitors of Cyclin-Dependent Kinase 2: Synthesis, X-Crystallographic Analysis, and Biological Activities", Journal of Medical Chemistry, Page EST:22.3, A-W.										
		AR	Mettey et al., "Aloisines, a New Family of CDK/GSK-3 Inhibitors. SAR Study, Crystal Structur in Complex with CDK2, Enzyme Selectivity, and Cellular Effects", J. Med. Chem. (2003), 46(2): 222-236.								
-	\vdash	AS	Novinson et al., "Synthesis and Antifungal Properties of Certain 7-Alkylaminopyrazolo[1,5-a]pyrimidines", J. Med. Chem. (1977), 20(2): 296-299.								
		AT	Senderowicz et al., "Phase I Trial of Continuous Infusion Flavopiridol, a Novel Cyclin-Dependent								
			Kinase Inhibitor, in Patients with Refractory Neoplasms", Journal of Clinical Oncology (Septemb 1998), 16(9): 2986-2999.								
		AU	the Cyclin-Depender	er et al., "Blochemical and Cellular Effects of Roscovitine, a Poten Cyclin-Dependent Kinases CDC2, CDK2 and CDK5", Eur. J. Bioch				hem.	(1997),	243:527	-536.
·	AV Bible et al., "Cytotoxic Synergy Between Flavopiridol (NSC 649890, L86-8275) and Various							August			
		AW	Shiota et al., "Synthesis and Structure-Activity Relationship of a New Series of Potent Angiotensin II Receptor Antagonists: Pyrazolo[1,5-a]pyrimidine Derivatives", Chem. Pharm. Bull. (1999), 47(7): 928-938.								
AX Yasuo Makisumi, "Studies on the Azaindolizine Compounds. XI. Synthesis of 6,7-Disub Pyrazolo[1,5-a]pyrimidines.", Chem. Pharm. Bull (1982), 19: 620-626/						isubstitu	ited				
EXAMINER 10 1 DATE CONSIDERED											
	. /	M		L.B.		. //	16	$\overline{}$			

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.